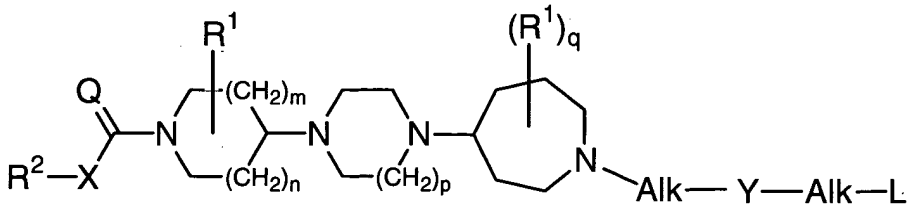
Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (Original) A compound according to the general Formula (I)



(I)

the pharmaceutically acceptable acid or base addition salts thereof, the stereochemically isomeric forms thereof, the *N*-oxide form thereof and prodrugs thereof, wherein :

n is an integer, equal to 0, 1 or 2;

m is an integer, equal to 1 or 2, provided that if m is 2, then n is 1;

p is an integer equal to 1 or 2;

q is an integer equal to 0 or 1;

Q is O or NR^3;

X is a covalent bond or a bivalent radical of formula -O-, -S- or $-NR^3-$;

each R^3 independently from each other, is hydrogen or alkyl;

each R^1 independently from each other, is selected from the group of Ar^1, Ar^1-alkyl and di(Ar^1)-alkyl;

R^2 is Ar^2, Ar^2-alkyl, di(Ar^2)alkyl, Het^1 or Het^1-alkyl;

Y is a covalent bond or a bivalent radical of formula -C(=O)-, $-SO_2-$ >C=CH-R or >C=N-R, wherein R is H , CN or nitro ;

each Alk represents, independently from each other, a covalent bond; a bivalent straight or branched, saturated or unsaturated hydrocarbon radical

having from 1 to 6 carbon atoms; or a cyclic saturated or unsaturated hydrocarbon radical having from 3 to 6 carbon atoms; each radical optionally substituted on one or more carbon atoms with one or more alkyl, phenyl, halo, cyano, hydroxy, formyl and amino radicals;

L is selected from the group of hydrogen, alkyl, alkyloxy, Ar^3-oxy, alkyloxycarbonyl, mono- and di(alkyl)amino, mono-and di(Ar^3)amino, Ar^3, Ar^3carbonyl, Het^2 and Het^2carbonyl;

Ar^1 is phenyl, optionally substituted with 1, 2 or 3 substituents, each independently from each other, selected from the group of halo, alkyl, cyano, aminocarbonyl and alkyloxy;

Ar^2 is naphthalenyl or phenyl, each optionally substituted with 1, 2 or 3 substituents, each independently from each other, selected from the group of halo, nitro, amino, mono- and di(alkyl)amino, cyano, alkyl, hydroxy, alkyloxy, carboxyl, alkyloxycarbonyl, aminocarbonyl and mono- and di(alkyl)aminocarbonyl;

Ar^3 is naphthalenyl or phenyl, optionally substituted with 1, 2 or 3 substituents, each independently from each other, selected from the group of alkyloxy, alkyl, halo, hydroxy, pyridinyl, morpholinyl, pyrrolidinyl, imidazo[1,2-*a*]pyridinyl, morpholinylcarbonyl, pyrrolidinylcarbonyl, amino and cyano;

Het^1 is a monocyclic heterocyclic radical selected from the the group of pyrrolyl, pyrazolyl, imidazolyl, furanyl, thienyl, oxazolyl, isoxazolyl, thiazolyl, isothiazolyl, pyridinyl, pyrimidinyl, pyrazinyl and pyridazinyl; or a bicyclic heterocyclic radical selected from the group of quinolinyl, quinoxalinyl, indolyl, benzimidazolyl, benzoxazolyl, benzisoxazolyl, benzothiazolyl,

benzisothiazolyl, benzofuranyl and benzothienyl; each heterocyclic radical may optionally be substituted on any atom by a radical selected from the group of halo and alkyl;

Het^2 is a monocyclic heterocyclic radical selected from the group of pyrrolidinyl, dioxolyl, imidazolidinyl, pyrrazolidinyl, piperidinyl, morpholinyl, dithianyl, thiomorpholinyl, piperazinyl, imidazolidinyl, tetrahydrofuranyl, 2H-pyrrolyl, pyrrolinyl, imidazolinyl, pyrrazolinyl, pyrrolyl, imidazolyl, pyrazolyl, triazolyl, furanyl, thienyl, oxazolyl, isoxazolyl, thiazolyl, thiadiazolyl, isothiazolyl, pyridinyl, pyrimidinyl, pyrazinyl, pyridazinyl and triazinyl;

or a bicyclic heterocyclic radical selected from the group of benzopiperidinyl, quinolinyl, quinoxalinyl, indolyl, isoindolyl, chromenyl, benzimidazolyl, imidazo[1,2-*a*]pyridinyl, benzoxazolyl, benzisoxazolyl, benzothiazolyl, benzisothiazolyl, benzofuranyl and benzothienyl;

each radical optionally substituted with one or more radicals selected from the group of Ar^1, Ar^1alkyl, halo, hydroxy, alkyl, piperidinyl, pyrrolyl, thienyl, oxo, alkyloxy, alkyloxyalkyl and alkyloxycarbonyl; and

alkyl is a straight or branched saturated hydrocarbon radical having from 1 to 6 carbon atoms or a cyclic saturated hydrocarbon radicals having from 3 to 6 carbon atoms; optionally substituted on one or more carbon atoms with one or more radicals selected from the group of phenyl, halo, cyano, oxo, hydroxy, formyl and amino.

2. (Original) A compound according to claim 1, characterized in that

n is 1;

m is 1;
p is 1;
q is 0;
Q is O;
X is a covalent bond;
each R^1 is Ar^1 or Ar^1-alkyl;

R^2 is Ar^2;
Y is a covalent bond or a bivalent radical of formula -C(=O)- ;
each Alk represents, independently from each other, a covalent bond
L is selected from the group of hydrogen, alkyloxy, Ar^3 and Het^2;
Ar^1 is phenyl;
Ar^2 is phenyl, optionally substituted with 1, 2 or 3 alkyl radicals;
Ar^3 is phenyl, optionally substituted with 1, 2 or 3 substituents, each independently from each other, selected from the group of alkyl and halo;
Het^2 is a monocyclic heterocyclic radical selected from the group of pyrazolyl, furanyl and isoxazolyl, each radical optionally substituted with one or more alkyl radicals; and
alkyl is a straight hydrocarbon radical having 1 to 6 carbon atoms, optionally substituted with one or more halo radicals.

3. (Currently Amended) A compound according to Claim 1 ~~any of claims 1-2, characterized in that~~ wherein R^1 is Ar^1methyl and attached to the 2-position or R^1 is Ar^1 and attached to the 3-position.

4. (Currently Amended) A compound according to ~~any of claims 1-3, characterized in that~~ Claim 1 wherein the R^2-X-C(=Q)- moiety is 3,5-di-(trifluoromethyl) phenylcarbonyl.

5. (Currently Amended) A compound according to ~~any of claims 1-4, characterized in that~~ Claim 1 wherein p is 1.

6. (Currently Amended) A compound according to ~~any of claims 1-5, characterized in that~~ Claim 1 wherein Y is -C(=O)-.

7. (Currently Amended) A compound according to ~~any of claims 1-6, characterized in that~~ Claim 1 wherein Alk is a covalent bond.

8. (Currently Amended) A compound according to ~~any of claims 1-3, characterized in that~~ Claim 1 wherein L is Het^2.

9. (Canceled)

~~11.~~ 10. (Currently Amended) A compound according to ~~any one of claims 1-10~~ claim 1 for use as an orally active~~, central penetrating~~ medicine.

~~12.~~ 11. (Currently Amended) ~~The use of a compound according to any one of claims 11 for the manufacture of a medicament for treating~~ A method for the treatment and/or prophylaxis of tachykinin mediated conditions comprising administering to a human in need of such administration of an effective amount of a compound according to claim 1.

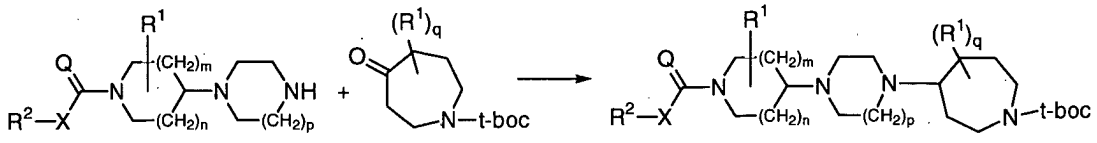
~~13.~~ 12. (Currently Amended) ~~The use of a compound according to claim 1-11 for the manufacture of a medicament for treating~~ A method for the treatment and/or prophylaxis of schizophrenia, emesis, anxiety, depression, irritable bowel syndrome (IBS), circadian rhythm disturbances, pain, neurogenic inflammation, asthma,

micturition disorders such as urinary incontinence and nociception comprising administering to a human in need of such administration of an effective amount of a compound according to claim 1.

~~14.~~ 13. (Currently Amended)A pharmaceutical composition comprising a pharmaceutically acceptable carrier and, as active ingredient, a therapeutically effective amount of a compound according to ~~any one of claims 1-9~~ claim 1.

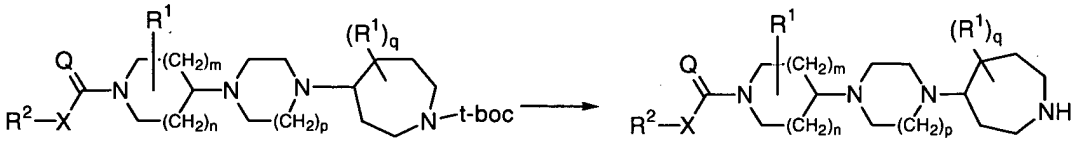
~~15.~~ 14. (Currently Amended)A process for preparing a pharmaceutical composition ~~as claimed in claim 14, characterized in that a pharmaceutically~~ comprising mixing a pharmaceutically acceptable carrier ~~is intimately mixed~~ with a therapeutically effective amount of a compound ~~as claimed in any one~~ of ~~claims 1-9~~ Claim 1.

~~16.~~ 15. (Currently Amended) A process for the preparation of a compound of Formula (I'') in which an intermediate compound of Formula (II) is reacted with an intermediate compound of Formula (III), wherein the radicals R^2, X, Q, R^1, m, n, p and q are as defined in claim 1.



(II) (III) (I'')

~~17.~~ 16. (Currently Amended) A process for the preparation of a compound of Formula (I') in which a final compound of Formula (I'') is reductively hydrogenated, wherein the radicals R^2, X, Q, R^1, m, n, p and q are as defined in claim 1.



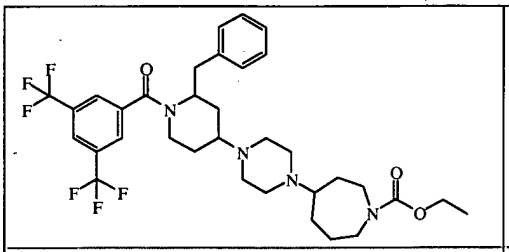
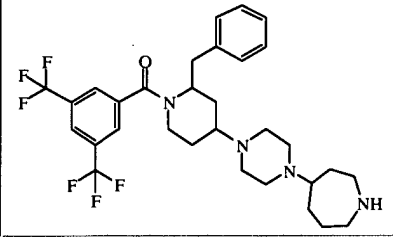
(I'') (I')

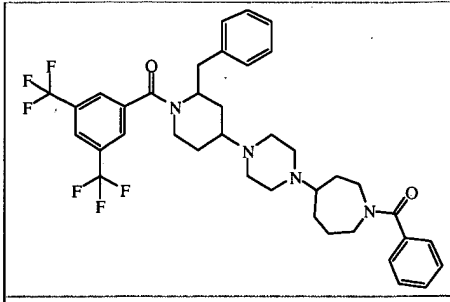
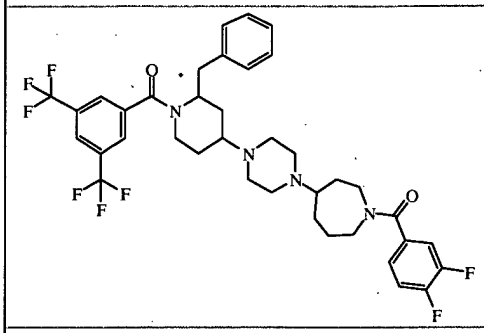
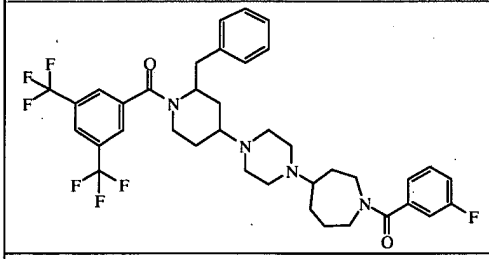
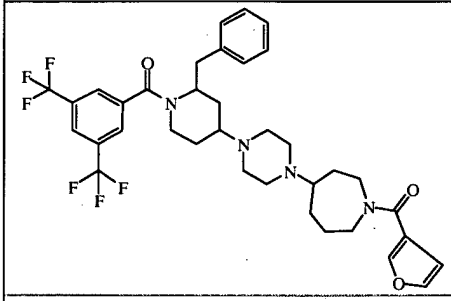
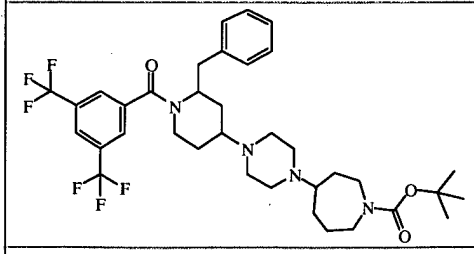
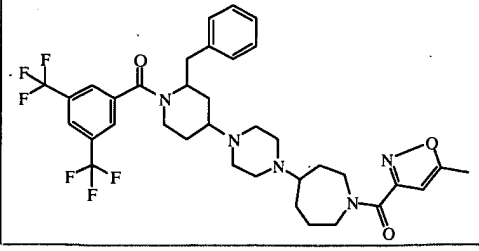
~~18.~~ 17. (Currently Amended) A process for the preparation of a compound according to Formula (I') comprising the consecutive steps of

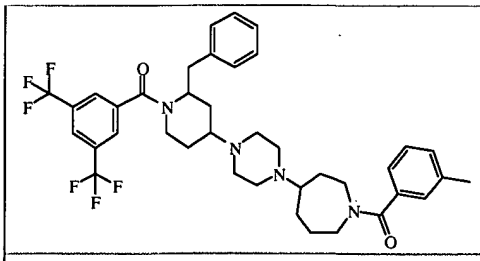
1) obtaining a compound of Formula (I'') according to claim ~~16~~ 15;

2) obtaining a compound of Formula (I') according to claim ~~17~~ 16.

18. (New) A compound select from the group consisting of

	2R-trans
	2R-trans

	2R-trans
	2R-trans
	2R-trans
	2R-trans
	2R-trans
	2R-trans

	2R-trans
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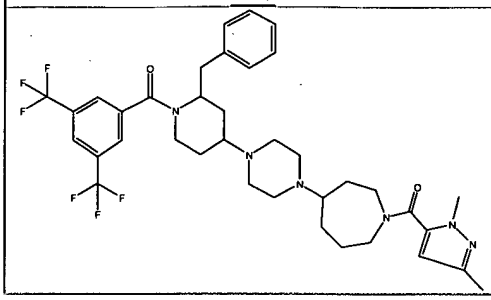	2R-trans